

U.S. DEPARTMENT OF LABOR WORKPLACE STANDARDS ADMINISTRATION BUREAU OF LABOR STANDARDS

MATERIAL SAFETY DATA SHEET

FORM NO OSHA-20 (MODIFIED) MAY 1971

MDC CONTROL NO. 1924

SECTI	ON I: MATE	RIAL AND MA	NUFACTURER IDENTIFICATION			
MANUFACTURER'S NAME EMERGENCY TELEPHO						
Idylite Company - Division of Ox ADDRESS (NUMBER, STREET, CITY, STATE AND Z				262-4101		<u>, , , , , , , , , , , , , , , , , , , </u>
3628 E. Olympic Blvd., Los Angel CHEMICAL NAME AND SYNONYMS	es, Cal	ifornia 9	TRADE NAME AND SYN	ONYMS	-	
Electropolishing Solution			Battelle #30			
CHEMICAL FAMILY	 		FORMULA			- Internation
Inorganic Acid Mixture			Proprietary			
	SECT	ON II: HAZAR	DOUS INGREDIENTS*			
PAINTS, PRESERVATIVES/SOLVENTS	%	TLV (UNITS)	ALLOYS AND METALL	%	ŢĻŲ (UNITS)	
PIGMENTS			BASE METAL			
CATALYST			ALLOYS			
VEHICLE		-	METALLIC COATINGS			
SOLVENTS			FILLER METAL PLUS COATING OR CORE FLUX			
ADDITIVES		,	OTHERS			
OTHERS	_				1	
HAZARDOUS	MIXTURE	S OF OTHER I	LIQUIDS, SOLIDS, OR GASES*	:	%	TLV (UNITS)
Chromic Acid (CrO ₂)	_				20	TFC in O.11 Mg/m
Phosphoric Acid (H_PO_1)					80	TXC in 100 Mg/m
		COTION III.	NIVALA LA PARA			
 		SECTION III: P	HYSICAL DATA SPECIFIC GRAVITY (H ₂ O = 1)			
BOILING POINT (OF)		 360°F	SPECIFIC GRAVITY (H ₂ U = 1)		1.7-1.8	
		700 1	PERCENT VOLATILE			NONE
VAPOR PRESSURE (mm Hg.)		BY VOLUME (%)			1401413	
VAPOR DENSITY (AIR = 1)			EVAPORATION RATE (= 1)			
SOLUBILITY IN WATER		100%			_	
APPEARANCE AND ODOR dark red-brown	. viscos	e. soluti	on, slight acid odor			
	SECTION IV	: FIRE AND E	XPLOSION HAZARD DATA			
FLASH POINT (METHOD USED) NONE		·	FLAMMABLE LIMIT	S L	91 1.	Uel
EXTINGUISHING MEDIA Water					·	
SPECIAL FIRE FIGHTING PROCEDURES NONE, use	copius	amounts o	of water			
		. ,				
UNUSUAL FIRE AND EXPLOSION HAZARDS						

				SF	CTIO	N V HFAI	LTH HAZA	RD DAT	Δ	
THRESHOLD LIMIT VALUE					<u> </u>	V V - 112.11	_ 111 11/12/	IND DI		
THE OF OVEREY POO		Section	II							
EFFECTS OF OVEREXPOS		ge to s	kin	and i	muco	ous mer	nbranes	:		
	Dalila	a or agr	IV.TII	and		Jub mer				
						<u>.</u> .				
EMERGENCY AND FIRST A										
	<u>Exte</u>	ernal -	TLUS	sh Wl	tn i	water				
	Inte	ernal -	give	e lar	ge (quanti	ties of	`wate	r then milk of magnesia - report to	
		-	doct	tor				_		
			uoo		F O TI	N 71. 55	. A O. T. I. U. T. I.			
	'		Ť			ON VI: KE S TO A VO	ACTIVITY	DATA	 	
0.T. D.U. I.T.	UNSTABLE		John Hong to Avois							
STABILITY	SI	TABLE	XX							
INCOMPATIBILITY (MATER										
Alkalis, reducing			rgai	nics						
HAZARDOUS DECOMPOSIT	ION PROD	UCTS							· · · · · · · · · · · · · · · · · · ·	
Toxic chromium p	roducts	if exp	ose	d to	fire					
HAZARDOUS		MAY OCC	UR			CONDIT	TONS TO A	VOID		
POLYMERIZATION		WILL NO.			+	 			<u> </u>	
		WILL NOT		UK	XX	<u> </u>	_			
								,		
				SECTIO	N VI	I: SPILL (OR LEAK I	PROCEDI	IIRES	
STEPS TO BE TAKEN IN C				ED OR	SPILL	_ED		INDULD	SNE0	
Flush with water	, neutr	ralize w	rith	sodi	um (carbona	ate.			
*		<u> </u>		-		· · · · · · · · · · · · · · · · · · ·				
W									···	
WASTE DISPOSAL METHOD Dilute with water	, r. redu	ıce hexi	val	ent c	hror	me wit	h hydro	sulfi	te and neutralize with sodium	
carbonate or sod	ium hyd	droxide.								
							_	•		
			SE U	י אחודי	111. CI	DECIAL D	ROTECTIO	א ואבטו	DMA TION	
RESPIRATORY PROTECTION	ON (SPECI	FY TYPE)	-							
exhaust ventilat			nly	when	. so	Lution	is ele	ctrif		
	LOCAL EXHAUST A tank while operating							SPECIAL scrub fumes for acid and chrome		
VENTILATION	MECHANICAL (GENERAL)								OTHER	
		COMMONE (GENERAL)								
PROTECTIVE GLOVES								ROTECTI		
yes, neoprene OTHER PROTECTIVE EQU	IDMENIT						<u>у</u> е	es, go	ggles	
apron, neoprene	ILMENI									
	-			SEC	TION	IX: SPEC	IAL PREC	AUTIONS	\$	
PRECAUTIONS TO BE TAKE Do not store in	EN IN HAI excess	NDLING AN heat or	D STO	RING					r reducing agents. Carboys should	
		-							· · · · · · · · · · · · · · · · · · ·	
be vented period	icly to	releas	se p	ressu	re.	Vent	more i	reque	ently in warm weather.	
OTHER PRECAUTIONS										
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Clair Mi.	ton	ul)	/ 			<u>Augu</u>	st 2,	L972		
PREPARED BY						DAT				

BATTELLE #30

SUPERSURFACING SOLUTION FOR STEEL

THE

UDYLITE GONPONNION

3523 E. GLYMPIG BLVD. • LGS MISBELS, EMLE, 83323

World's Largest Plating Supplier

INTRODUCTION

The supersurfaced metal has distinct property advantages and is an improvement over the abrasive-finished or machine-cut surface in several important aspects. The supersurface has:

- rreedom from mechanically damaged surface metal, stresses, slivers, and "debris" introduced by the cutting, tearing, working, etc., caused by abrasive finishing, lathe turning, grinding, broaching, etc. There is no "weak metal skin" to suffer fatigue and spall off slivers and flakes of metal, or to pit, etc., in engineering component service.
- b. Freedom from the directional effect of mechanical finishing methods. (Tumbling, of course, reveals no directional effect, but is rarely used for high-quality surface finishing of shafts, pins, gear teeth, etc.
- c. Easily discernable defects, so that failure of a part is forestalled, whereas abrasive finishing actually covers over defects.

APPLICATIONS FOR SUPERSURFACING

The principal applications are for engine and other mechanical parts made of expensive, special alloy steels. These alloys when fabricated require the removal of the mechanically damaged surface skin. If this is not removed the peak performance or the best wearing quality of the alloy cannot be realized. Also, it is for those parts where surface smoothnesses should be finer than 5 r.m.s. microinches, down to as fine as 0.1 microinch.

To achieve the best results by supersurfacing, the surface must be brought first to a good quality finish by mechanical means; to about 7 to 10 r.m.s. microinch finish. The steel must be relatively clean and free from local stresses, such as those introduced by careless grinding into very hard steels. Removal of the outer most surface might release stresses and produce cracking.

COMPARISON OF SUPERSURFACING TO ELECTROPOLISHING

For many applications, electropolishing in the #31 bath, as described in the bulletin on steel, will produce entirely satisfactory results. Thus, all points, except 3 and 4 in the following, direct first consideration to the #31 bath if it can possibly do the job.

- a. The supersurfacing bath has a finite life and must be operated on a decant schedule along lines discussed in the stainless steel electropolishing bulletin.
- b. Operating cost is higher for the supersurfacing bath; about 4 to 5 times the cost of electropolishing steels in the #31 bath (about 2 to 4 times the cost of electropolishing stainless steels).
- c. The supersurface is much more rust resistant than the electropolished in the #31 bath.
- d. Supersurfacing gives better results with steels that

tend to be dirty or to have duplex phase structures.

e. Supersurfacing requires a shorter time, 30-90 seconds, and much higher current densities (and tank voltages) 12 to 24 amps per square inch.

Contamination control of the supersurfacing bath is more critical than that of the #31 bath.

PREPARATION AND MAINTENANCE OF THE SUPERSURFACING BATH

Operation

The working temperature is between 60 and 100 C. (140° to 212°F.).

Current density can vary over a wide range, depending on the particular application, and C.D.'s of 200 to 2000 amperes per square foot may be used. Suitable current densities are given below:

Deburring

1,000 - 2,000 amperes per square foot

1,000 amperes per square foot

Descaling

250 - 500 amperes per square foot

The specific gravity of a new solution is 1.73 at 70°F. The specific gravity of a used solution is from 1.79 to 1.80 at temperatures between 70° and 75°F. Solutions above this figure should be decanted and fresh solution added.

The following is the maximum amount of dissolved solids:

Aluminum 0.1% by weight Iron 2.7% by weight Trivalent Chromic Oxide 3.0% by weight

The time of the treatment depends upon the amount of metal to be removed. The amount of metal removed is also influenced by the composition and condition of the steel being treated, and, therefore, it is necessary to check the loss experimentally where presion is required. When the dissolution rate is established for a particular C.D. and racking arrangement, it is possible to reproduce results accurately. The distribution of metal removal is, of course, influenced by the position of cathodes and/or shields, and these should be constructed and arranged with precision. Components should also be placed in the solution so that the form offers the least resistance to the flow of bath by stirring. It is important that the surfaces undergoing treatment should make the fullest possible contact with the rapidly moving liquid. Auxiliary cathodes may be aluminum or copper, particularly copper where absence of space demands a conductor with the lowest sectional resistance. Shields may be made of Bakelite or P.V.D. The protection of edges may be ensured with auxiliary anodes made of steel or aluminum, but not copper, as the anodes are attacked.

The purpose of the rapid stirring by stainless steel (Type 317) propellers is to rapidly remove the heat produced at the anode, and which, apart from spoiling the polishing effect, also decomposes chromic acid which is wasteful.

The parts are to be held on steel or aluminum racks. Copper and brass are unsuitable. The high current densities necessitate the use of stout racks and electrical contact. Where posible, contact should be made with any part of the component which is not immersed. Tapped holes or screw clamps are recommended to ensure

good contact. Spring clips and magnetic jigs may also be used. In repetition work, the rack, apart from the contact surface, should be protected with P.V.C. tape, or a plastisol. Failure of the coatings is often due to faulty rack design and results in overheating.

When the iron content reaches the maximum value, further work is impossible. The iron content may be reduced by discarding a portion of the bath and adding a sufficient quantity of new solution make up the required volume.

Steel racks dissolve at about the same rate as the work and must be replaced as needed. Aluminum racks dissolve at a relatively less rate, as shown by the following table, and thus have better life than steel racks. The table shows loss on an aluminum rack to be about 1/2 the area of the steel part. The aluminum-to-steel loss ratio changes slightly as the bath becomes aged. The operating level at 2.4 to 2.7 per cent dissolved iron is the one to maintain, by discarding a small amount of used bath at 2.7 per cent iron.

RATIO OF ALUMINUM RACK LOSS TO STEEL PART LOSS IN SUPERSURFACING VS. USE OF THE BATH

Specific	Age	Wei	ight Loss,	% Fe		
Gravity	ravity amp-hr/l		Steel	Steel/Al	in Bath	Volts
1.70	0	0.031	0.325	10	0 <u></u> .	6.35
1.715	53	0.021	0.262	12.4	0.71	7.1
1.73	100	0.023	0.217	9.5	1.52	7.7
1.75	153	0.0301	0.183	6	2,21	9,5
.1.76	192*	0.0264	0.144	5.5	2.71	9.8
1.715	_. 192	0.039	0.186	4.8	2.89	9,1

^{*} Amp-hr/gal. = 730

Precautions

Solution is lost by dragout, and components which have been treated should be well drained before rinsing to conserve the acids. The steel, alloy steel, or aluminum components are not attacked chemically while they are immersed in the solution.

To ensure that chromic acid is not reduced unnecessarily, it is important that, in addition to the avoidance of overheating mentioned previously, grease, copper and copper alloys should not be allowed to enter the bath. Sulphuric acid can also cause the reduction of chromic acid during the electropolishing of steel and, consequently, if a pretreatment bath containing sulphuric acid is used, thorough rinsing should be carried out before supersurfacing.

A high working temperature is necessary to ensure good agitaion, a low voltage, and the retention of chromic acid in solution. A low specific gravity will cause loss of throwing power and a nonuniform removal of the metal.

Staining may be caused by a high iron content or overheating. The latter may be caused by poor contacts, faulty agitation, or too long a period of immersion. It is sometimes better to give a number

of short treatments rather than a continuous treatment when heavy metal removal is required.

Persistant defects under satisfactory conditions may be associated with the material under treatment. Variations in the amount of cold deformation may cause a variation in smoothness, due to variations in structure. This effect is also noticeable on components machined from bar stocks and sections where the original surface possesses different properties from the core material exposed during machining.

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Los Angeles